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10	BEFORE THE MEDICAL BOARD OF CALIFORNIA
11	DEPARTMENT OF CONSUMER AFFAIRS STATE OF CALIFORNIA
12	STATE OF CALAFORNA
13	In the Matter of the First Amended Accusation   Case No. 800-2020-067207
14	Against:  FIRST AMENDED ACCUSATION
15	SARAH ASH COMBS, M.D. 3700 10th Avenue, Apt. 3H
16	San Diego, CA 92103
17	Physician's and Surgeon's Certificate No. A 125860,
18	Respondent.
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20	Complainant alleges:
21	<u>PARTIES</u>
22	1. William Prasifka (Complainant) brings this First Amended Accusation solely in his
23	official capacity as the Executive Director of the Medical Board of California, Department of
24	Consumer Affairs (Board).
25	2. On or about May 22, 2013, the Medical Board issued Physician's and Surgeon's
26	Certificate No. A 125860 to Sarah Ash Combs, M.D. (Respondent). The Physician's and
27	Surgeon's Certificate expired on December 31, 2016, and has not been renewed.
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(c) Repeated negligent acts. To be repeated, there must be two or more negligent acts or omissions. An initial negligent act or omission followed by a separate and distinct departure from the applicable standard of care shall constitute repeated negligent acts.

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# FIRST CAUSE FOR DISCIPLINE

# (Repeated Negligent Acts)

- 10. Respondent has subjected her Physician's and Surgeon's Certificate No. A 125860 to disciplinary action under sections 2227 and 2234, as defined by section 2234, subdivision (c), of the Code, in that she committed repeated negligent acts in her care and treatment of Patient A, as more particularly alleged hereinafter:<sup>1</sup>
- 11. On or about the night of January 6, 2016, Patient A, who was a teenager, presented at the emergency department of Rady Children's Hospital in San Diego, California. According to Patient A's father, Patient A had intentionally ingested seven tablets of Midol and 10 tablets of iron approximately one and a half hours prior to arrival.
- 12. At the time, Respondent was a board-certified pediatrician and in training for her subspecialty fellowship in pediatric emergency medicine. Respondent provided care and treatment to Patient A, alongside Dr. S.L., her direct supervisor.
- 13. Upon arrival at the emergency department, Patient A complained of dizziness, nausea, and vomiting. Patient A was ordered ondansetron for her dizziness and nausea at approximately 23:39, followed by a fluid bolus approximately two and a half hours later.
- 14. Patient A's vital signs were taken, showing a heart rate of 133 beats per minute, which was markedly tachycardic. An EKG also showed a heart rate of 108 beats per minute. However, Patient A's cardiovascular status was noted on the physical exam as follows: "Normal rate, regular rhythm and normal heart sounds."
- 15. According to the ED Provider Notes, lab tests were to be ordered, including an iron level test. However, on or about January 7, 2016, at approximately 01:55, a ferritin level test was erroneously ordered, not an iron level test.<sup>2</sup> The lab results reported the findings for "Ferritin," which were received at approximately 02:57. According to the lab results, the ferritin test showed

<sup>&</sup>lt;sup>1</sup> References to "Patient A" herein are used to protect patient privacy.

<sup>&</sup>lt;sup>2</sup> Ferritin is a protein that stores iron inside the cells. A ferritin test measures the level of ferritin in the body. Ferritin levels indicate the amount of stored iron, but they do not measure the iron outside of the cells. An iron test, in contrast, measures the amount of iron in the blood. After a suspected overdose of iron, a serum iron level is the most appropriate test to order to assess for acute toxicity.

a level of "8" ng/mL, with a reference range of 6-70 ng/mL. There were no lab results for iron. Nevertheless, the ED Provider Notes stated: "Labs as per below, grossly WNL... Iron well below threshold." The Plan and Discharge Instructions further stated: "Your iron level here was normal."

- 16. At approximately 04:00 on or about January 7, 2016, Patient A complained of additional nausea to the ED nurse. At approximately 04:02, an additional dose of ondansetron was ordered, which was administered at 04:07. Approximately one hour later, Patient A was discharged home, with the last physician re-assessment occurring at approximately 04:12. Prior to discharge, there was no assessment done and no documentation made as to the etiology of Patient A's continuing nausea and whether or not it would persist.
- 17. Following discharge, Patient A subsequently developed severe abdominal and chest pain and returned to the emergency department later the same day. Patient A's lab results showed a hemoglobin of 11.1 g/dL. Patient A was found to be in fulminant liver failure due to iron overdose and required an emergency liver transplant.
- 18. On or about January 14, 2021, Respondent was interviewed in connection with the Board's investigation regarding her care and treatment of Patient A. Respondent stated that she intended to order a total body iron test for Patient A, not a ferritin test. When placing the order, Respondent explained that she typed the word "iron" into the electronic medical record system. According to Respondent, the system automatically defaulted to "ferritin" and, as a result, a ferritin test was ordered instead of an iron test. However, the lab test options that appeared on the screen were actually as follows, from top to bottom:

### FERRITIN (IRON)

#### **IRON**

## IRON + TIBC

Despite "IRON" appearing on the screen, the test for iron was not ordered.

19. At her Board interview, Respondent further stated that when she reviewed the labs, the results appeared as either "ferritin paren iron" or "iron paren ferritin." Despite the word "ferritin" appearing on the results, Respondent stated she expected that the results were

measuring what she thought she had ordered (*i.e.*, an iron level), so she "glanced past it, as you often do in the emergency room, [and] saw a normal level . . . ." According to Respondent, she presumed that the result would likely be normal given the amount of iron tablets that Patient A had reportedly ingested.

- 20. Respondent committed repeated negligent acts in her care and treatment of Patient A, which included, but were not limited to the following:
  - (i) Respondent failed to order the correct test to assess for acute iron toxicity and she failed to appropriately review and interpret the test results received;
  - (ii) Respondent failed to properly assess and document the etiology of Patient A's continuing nausea and whether or not the patient's nausea would persist; and
    - (iii) Respondent failed to properly document Patient A's tachycardia.

# SECOND CAUSE FOR DISCIPLINE

# (Failure to Maintain Adequate and Accurate Medical Records)

21. Respondent has subjected her Physician's and Surgeon's Certificate No. A 125860 to disciplinary action under sections 2227 and 2234, as defined by section 2266, of the Code, in that she failed to maintain adequate and accurate records regarding her care and treatment of Patient A, as more particularly alleged in paragraphs 10 through 20, above, which are hereby incorporated by reference and re-alleged as if fully set forth herein.

#### PRAYER

WHEREFORE, Complainant requests that a hearing be held on the matters herein alleged, and that following the hearing, the Medical Board of California issue a decision:

- 1. Revoking or suspending Physician's and Surgeon's Certificate No. A 125860, issued to Respondent Sarah Ash Combs, M.D.;
- 2. Revoking, suspending or denying approval of Respondent Sarah Ash Combs, M.D.'s authority to supervise physician assistants, pursuant to section 3527 of the Code, and advanced practice nurses;

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